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INFORMATION DISCLOSURE
STATEMENT BY APPLICANT

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Sheet 1 of 2

3226087

Complete if Known

Application Number	10/602,693
Filing Date	June 20, 2003
First Named Inventor	Sommadossi et al.
Group Art Unit	1623
Examiner Name	Travis C. McIntosh, III
Attorney Docket Number	06171.105070 IDX 1006 DIV1 US

U.S. PATENT DOCUMENTS

Examiner Initials *	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear	T ⁶
		Number	Kind Code (if known)				
(7)	AA	6,455,508	B1	Ramasamy et al.	09-24-2002		
	AB	6,495,677	B1	Ramasamy et al.	12-17-2002		
	AC	6,566,344	B1	Gosselin et al.	05-20-2003		
	AD	6,566,365	B1	Storer	05-20-2003		
	AE	6,569,837	B1	Gosselin et al.	05-27-2003		
	AF	6,605,614	B2	Bachand et al.	08-12-2003		
	AG	6,660,721	B2	Devos et al.	12-09-2003		
	AH	6,748,161	B2	Ko et al.	06-08-2004		
	AJ	6,777,395	B2	Bhat et al.	08-17-2004		
	AJ	6,784,166	B2	Devos et al.	08-31-2004		
	AK	6,812,219	B2	LaColla et al.	11-02-2004		
	AL	6,815,542	B2	Hong et al.	11-09-2004		
	AM	6,831,069	B2	Tam, et al.	12-14-2005		
	AN	6,908,924	B2	Watanabe, et al.	06-21-2005		
	AO	6,911,424	B2	Schinazi, et al.	06-28-2005		
	AP	6,914,054	B2	LaColla et al.	07-05-2005		
	AQ	2002-0099072	A1	Bachand et al.	07-25-2002		
	AR	2002-0156030	A1	Ramasamy et al.	10-24-2002		
	AS	2003-0008841	A1	Devos et al.	01-09-2003		
	AT	2002-0147160	A1	Bhat, et al.	10-10-2002		
	AU	2003-0220290	A1	Gosselin et al.	11-27-2003		
	AV	2003-0225028	A1	Gosselin et al.	12-04-2003		
	AW	2003-0225037	A1	Storer	12-04-2003		
	AX	2003-0236216	A1	Devos, et al.	12-25-2003		
	AY	2004-0002476	A1	Stuyver, et al.	01-01-2004		
	AZ	2004-0002596	A1	Hong et al.	01-01-2004		
	AAA	2004-0023921	A1	Hong, et al.	02-05-2004		
	AAB	2004-0059104	A1	Cook et al.	03-25-2004		
	AAC	2004-0063622	A1	LaColla et al.	04-01-2004		
	AAD	2004-0063658	A1	Roberts et al.	04-01-2004		
	AAE	2004-0067901	A1	Bhat et al.	04-08-2004		
	AAF	2004-0072788	A1	Bhat et al.	04-15-2004		

Examiner Signature

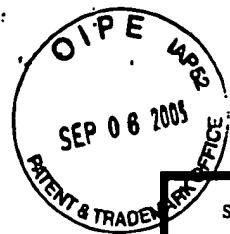
Date Considered

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Sheet 2 of 2

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Examiner Name	Travis C. McIntosh, III
Attorney Docket Number	06171.105070 IDX 1006 DIV 1 US

U.S. PATENT DOCUMENTS

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		Number	Kind Code (if known)				
(D)	BA	2004-0097461	A1	Sommadossi et al.	05-20-2004		
	BB	2004-0097462	A1	LaColla et al.	05-20-2004		
	BC	2004-0101535	A1	Sommadossi et al.	05-27-2004		
	BD	2004-0102414	A1	Sommadossi et al.	05-27-2004		
	BE	2004-0110717	A1	Carroll, et al.	06-10-2004		
	BF	2004-0110718	A1	Devos et al.	06-10-2004		
	BG	2004-0147464	A1	Roberts, et al.	07-29-2004		
	BH	2004-0248844	A1	Ismaili et al.	12-09-2004		
	BI	2005-0009737	A1	Clark, et al.	01-13-2005		
	BJ	2005-0090463	A1	Roberts, et al.	04-28-2005		
	BK	2005-0101550	A1	Roberts, et al.	05-12-2005		
	BL	2005-0107312	A1	Keicher, et al.	05-19-2005		
	BM	2005-0119200	A1	Roberts, et al.	06-02-2005		
	BN	2005-0124532	A1	Sommadossi et al.	06-09-2005		
	BO	2005-0137161	A1	Sommadossi et al.	06-23-2005		

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		Office ³	Number	Kind Code ² (if known)				
(D)	BP	FR	2,662,165	A1	Univ. Pier et Curie	11-22-1991	provided as Derwent Abstract	
(W)	BQ	JP	63-215694	A2	Yamasa Shoyu Co. Ltd.	09-08-1988	provided as Delphion Abstract	
(W)	BR	JP	06-228186	A2	Yamasa Shoyu Co. Ltd.	08-16-1994	provided as Delphion Abstract	

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁴

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				Filing Date	June 20, 2003
				First Named Inventor	Sommadossi et al.
				Group Art Unit	1614
				Examiner Name	Unassigned
Sheet	1	of	6	Attorney Docket Number	06171.105070 IDX 1006 DIV

3425621/1

U.S. PATENT DOCUMENTS

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		Number	Kind Code (if known)			
P	AA	3,480,613	A	Walton et al.	11-25-1969	
	AB	5,977,061	A	De Clercq	11-02-1999	
	AC	6,340,690	B1	Bachand et al.	01-22-2002	
	AD	6,348,587	B1	Schinazi et al.	02-2002	
	AE	6,395,716	B1	Gosselin et al. (Novirio / Idenix)	05-28-2002	
	AF	6,444,652	B1	Gosselin et al. (Novirio / Idenix)	09-03-2002	
	AG	6,573,248	B1	Ramasamy et al.	06-03-2003	
	AH	2002/0019363	A1	Ismaili et al.	02-2002	
	AI	2002/0055483	A1	Watanabe et al.	05-09-2002	
	AJ	2002/0147160	A1	Bhat et al.	10-10-2002	
	AK	2003/008841	A1	Devos et al.	01-09-2003	
	AL	2003/028013	A1	Wang et al.	02-06-2003	
	AM	2003/0050229	A1	Sommadossi et al.	03-13-2003	
	AN	2003/0060400	A1	LaColla et al.	03-27-2003	
	AO	2003/0083307	A1	Devos et al.	05-01-2003	
	AP	2003/0087873	A1	Stuyver et al.	05-08-2003	

FOREIGN PATENT DOCUMENTS

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		Office ³	Number	Kind Code ² (if known)				
P	AQ	FR	1,521,076	A	Merck & Co. Inc.	04-12-1968		
	AR	FR	1,581,628	A	Merck & Co. Inc.	09-19-1969		
	AS	FR	2,662,165	A	Univ. Paris Curie	11-22-1991		
	AT	GB	1,163,103	A	Merck & Co. Inc.	09-04-1969		
	AU	GB	1,209,654	A	Merck & Co. Inc.	10-21-1970		
	AV	JP	63-215694	A	Yamasa Shoyu Co. Ltd.	09-08-1988		
	AW	JP	06-228186	A	Yamasa Shoyu Co. Ltd.	08-16-1994		
	AX	WO	98/16184	A2	ICN Pharmaceuticals	04-23-1998		
	AY	WO	99/43691	A1	Emory U.; U.Ga.R.F.	02-09-1999		
	AZ	WO	00/09531	A2	Novirio Pharm. (Idenix)	02-24-2000		
	AAA	WO	01/32153	A2	Biochem Pharma	05-10-2001		

Examiner Signature		Date Considered	3/29/05
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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

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Sheet	2	of	6	Attorney Docket Number	06171.105070 IDX 1006 DIV
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FOREIGN PATENT DOCUMENTS

Examiner Initials	Cite No. ¹	Foreign Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁴
		Office ³	Number				
	BA	WO	01/60315	A2	Biochem Pharma	08-23-2001	
	BB	WO	01/68663	A1	ICN Pharmaceuticals	09-20-2001	
	BC	WO	01/79246	A2	Pharmasset	10-25-2001	
	BD	WO	01/90121	A2	Novirio Pharm. (Idenix)	11-29-2001	
	BE	WO	01/91737	A2	Novirio Pharm. (Idenix)	06-12-2001	
	BF	WO	01/92282	A2	Novirio Pharm. (Idenix)	06-12-2001	
	BG	WO	01/96353	A2	Novirio Pharm. (Idenix)	12-20-2001	
	BH	WO	02/03997	A1	ICN Pharmaceuticals	01-17-2002	
	BI	WO	02/18404	A2	F. Hoffmann-La Roche	03-07-2002	
	BJ	WO	02/32920	A2	Pharmasset	04-25-2002	
	BK	WO	02/48165	A2	Pharmasset	06-20-2002	
	BL	WO	02/057287	A2	Merck & Co. Inc.	07-25-2002	
	BM	WO	02/057425	A2	Merck & Co. Inc.	07-25-2002	
	BN	WO	02/070533	A2	Pharmasset	09-12-2002	
	BO	WO	02/094289	A1	F. Hoffmann-La Roche	11-28-2002	
	BP	WO	02/100415	A2	F. Hoffmann-La Roche	12-19-2002	
	BQ	WO	03/026589	A2	Idenix; CNRS; U. Montp.	04-03-2003	
	BR	WO	03/026675	A1	Idenix; CNRS; U. Montp.	04-03-2003	
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	BT	WO	03/061385	A1	Ribapharm	07-31-2003	
	BU	WO	03/061576	A2	Ribapharm	07-31-2003	
	BV	WO	03/062255	A2	Ribapharm	07-31-2003	
	BW	WO	03/062256	A1	Ribapharm	07-31-2003	
	BX	WO	03/062257	A1	Ribapharm	07-31-2003	
	BY	WO	03/063771	A2	Pharmasset	08-07-2003	
	BZ	WO	03/068162	A2	Pharmasset	08-21-2003	
	BAA	WO	03/072757	A2	Biota Inc.	09-04-2003	
V	BAB	WO	03/093290	A2	Genelabs Technologies	11-13-2003	
	BAC	WO	04/002422	A2	Idenix; Univ.D.S.Cagliari	01-08-2004	
	BAD	WO	04/002999	A2	Idenix; Univ.D.S.Cagliari	01-08-2004	

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

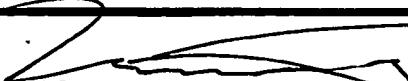
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				<i>Complete if Known</i>
				Application Number
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				Group Art Unit
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Sheet	3	of	6	06171.105070 IDX 1006 DIV

3425621 1

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
(L)	CA	ALTMANN <i>et al.</i> , "The synthesis of 1'-methyl carbocyclic thymidine and its effect on nucleic acid duplex stability," <i>Synlett, Thieme Verlag, Stuttgart, De</i> , 10:853-855 (1994).	
	CB	BAGINSKI, S. G, <i>et al.</i> , "Mechanism of action of a pestivirus antiviral compound," <i>PNAS USA</i> , 97(14):7981-7986 (2000).	
	CC	BEIGELMAN, L.N., <i>et al.</i> , "Epimerization during the acetolysis of 3-O-acetyl-5-O-benzoyl-1,2-O-isopropylidene-3-C-methyl- α ,D-ribofuranose. Synthesis of 3'-C-methylnucleosides with the β -D-ribo- and α -D-arabino configurations," <i>Carbohydrate Research</i> , 181:77-88 (1988).	
	CD	BEIGELMAN, L.N., <i>et al.</i> , "A general method for synthesis of 3'-C-alkylnucleosides," <i>Nucleic Acids Symp. Ser.</i> , 9:115-118 (1981).	
	CE	BERENGUER, M., <i>et al.</i> , "Hepatitis B and C viruses: Molecular identification and targeted antiviral therapies," <i>Proceedings of the Association of American Physicians</i> , 110(2), 98-112 (1998).	
	CF	CARROLL, S.S., <i>et al.</i> , "Inhibition of hepatitis C virus RNA replication by 2'-modified nucleoside analogs," <i>The Journal of Biological Chemistry</i> , 278(14):11979-11984 (2003).	
	CG	CZERNECKI, S., <i>et al.</i> , "Synthesis of various 3'-branched 2',3'-unsaturated pyrimidine nucleosides as potential anti-HIV agents," <i>J. Org. Chem.</i> , 57:7325-7328 (1992).	
	CH	De FRANCESCO, R., <i>et al.</i> , "Approaching a new era for hepatitis C virus therapy: Inhibitors of the NS3-4A serine protease and the NS5B RNA-dependent RNA polymerase," <i>Antiviral Research</i> , 58:1-16 (2003).	
	CI	FAIVRE-BUET, V., <i>et al.</i> , "Synthesis of 1'-deoxypsicofuranosyl-deoxynucleosides as potential anti-HIV agents," <i>Nucleosides & Nucleotides</i> , 11(7):1411-1424 (1992).	
	CJ	FARKAS, J., <i>et al.</i> , "Nucleic acid components and their analogues. XCIV. Synthesis of 6-amino-9-(1-deoxy- β -D-psicofuranosyl)purine", <i>Collect. Czech. Chem. Commun.</i> 32:2663-2667 (1967).	
	CK	FARKAS, J., <i>et al.</i> , "Nucleic acid components and their analogues. LXXIX. Synthesis of methyl 1-deoxy-D-psicofuranosides substituted at C ₍₁₎ with halo atoms or a mercapto group," <i>Collect. Czech. Chem. Commun.</i> , 31:1535-1543 (1996).	
	CL	FEDOROV, I.I., <i>et al.</i> , "3'-C-Branched 2'-deoxy-5-methyluridines: Synthesis, enzyme inhibition, and antiviral properties," <i>J. Med. Chem.</i> , 35(24):4567-4575 (1992).	
	CM	FRANCHETTI, P., <i>et al.</i> , "2'-C-Methyl analogues of selective adenosine receptor agonists: synthesis and binding studies," <i>J. Med. Chem.</i> , 41(10):1708-1715 (1998).	
V	CN	GROUILLER, A., <i>et al.</i> , "Novel p-toluenesulfonylation and thionocarbonylation of unprotected thymine nucleosides," <i>Synlett</i> , 1993, 221-222 (March 1993).	
(L)	CO	HARAGUCHI, K., <i>et al.</i> , "Preparation and reactions of 2'- and 3'- vinyl bromides of uracil nucleosides: Versatile synthons for anti-HIV agents," <i>Tetrahedron Letters</i> , 32(28):3391-3394 (1991).	

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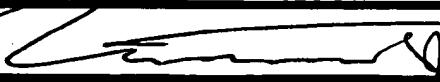
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				Group Art Unit	1614
				Examiner Name	Unassigned
Sheet	4	of	6	Attorney Docket Number	06171.105070 IDX 1006 DIV

3425621 1

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ^	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
<i>CW</i>	DA	HARAGUCHI, K., <i>et al.</i> , "Stereoselective synthesis of 1'-C-branched uracil nucleosides from uridine," <i>Nucleosides & Nucleotides</i> , 14(3-5):417-420 (1995).	
	DB	HARRY-O'KURU, R.E., <i>et al.</i> , "A short, flexible route toward 2'-C-branched ribonucleosides", <i>J.Org. Chem.</i> , 62:1754-1759 (1997). (Scheme 11).	
	DC	HARRY-O'KURU, R.E., <i>et al.</i> , "2'-C-Alkylribonucleosides: Design, synthesis, and conformation," <i>Nucleosides & Nucleotides</i> , 16(7-9):1457-1460 (1997). ("Rogers" in #2; correct name in #7)	
	DD	HATTORI, H., <i>et al.</i> , "Nucleosides and nucleotides. 175. Structural requirements of the sugar moiety for the antitumor activities of new nucleoside antimetabolites, 1-(3-C-ethynyl-b-D-ribo-pentofuranosyl)cytosine and -uracil," <i>J. Med. Chem.</i> , 41:2892-2902 (1998).	
	DE	HREBABECKY, H., <i>et al.</i> , "Nucleic acid components and their analogues. CXLIX. Synthesis of pyrimidine nucleosides derived from 1-deoxy-D-psicose," <i>Collect. Czech. Chem. Commun.</i> , 37:2059-2065 (1972).	
	DF	HREBABECKY, H., <i>et al.</i> , "Synthesis of 7- and 9β-D-psicofuranosylguanine and their 1'-deoxy derivatives" <i>Collect. Czech. Chem. Commun.</i> , 39:2115-2123 (1974).	
	DG	IINO, T., <i>et al.</i> , "Nucleosides and nucleotides. 139. Stereoselective synthesis of (2'S)-2'-C-alkyl-2'-deoxyuridines," <i>Nucleosides and Nucleotides</i> , 15(1-3):169-181 (1996).	
	DH	ITOH, Y., <i>et al.</i> , "Divergent and stereocontrolled approach to the synthesis of uracil nucleosides branched at the anomeric position," <i>J. Org. Chem.</i> , 60(3):656-662 (1995).	
	DI	JOHNSON, C.R., <i>et al.</i> , "3'-C-Trifluoromethyl ribonucleosides," <i>Nucleosides & Nucleotides</i> , 14(1&2):185-194 (1995).	
	DJ	KAWANA, M., <i>et al.</i> , "The deoxygenation of tosylated adenosine derivatives with Grignard reagents," <i>Nucleic Acids Symp. Ser.</i> , 17:37-40 (1986).	
	DK	LAVAIRE, S., <i>et al.</i> , "3'-Deoxy-3'-C-trifluoromethyl nucleosides: Synthesis and antiviral evaluation," <i>Nucleosides & Nucleotides</i> , 17(12):2267-2280 (1998).	
	DL	LEYSEN, P., <i>et al.</i> , "Perspectives for the treatment of infections with Flaviviridae," <i>Clinical Microbiology Reviews</i> (Washington, D.C.), 13(1):67-82 (January 2000).	
	DM	MARTIN, X., <i>et al.</i> , "Intramolecular hydrogen bonding in primary hydroxyl of thymine 1-(1-deoxy-β-D-psicofuranosyl) nucleoside," <i>Tetrahedron</i> , 50(22):6689-6694 (1994).	
<i>W</i>	DN	MATSUDA, A., <i>et al.</i> , "Radical deoxygenation of <i>tert</i> -alcohols in 2'-branched-chain sugar pyrimidine nucleosides: Synthesis and antileukemic activity of 2'-deoxy-2'(S)-methylcytidine," <i>Chem. Pharm. Bull.</i> , 35(9):3967-3970 (1987).	
<i>A</i>	DO	MATSUDA, A., <i>et al.</i> , "Alkyl addition reaction of pyrimidine 2'-ketonucleosides: Synthesis of 2'-branched-chain sugar pyrimidine nucleosides (Nucleosides and Nucleotides. LXXXI)," <i>Chem. Pharm. Bull.</i> , 36(3):945-953 (1988).	

Examiner Signature		Date Considered	3/29/05
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 5 of 6 Attorney Docket Number 06171.105070 IDX 1006 DIV				Complete If Known
				Application Number 10/602,693
				Filing Date June 20, 2003
				First Named Inventor Sommadossi et al.
				Group Art Unit 1614
				Examiner Name Unassigned

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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. ^{T²}
(a)	EA	MATSUDA, A., et al., "Nucleosides and Nucleotides. 94. Radical deoxygenation of <i>tert</i> -alcohols in 1-(2-C-alkylpentofuranosyl)pyrimidines: Synthesis of (2'S)-2'-deoxy-2'-C-methylcytidine, an antileukemic nucleoside," <i>J. Med. Chem.</i> , 34:234-239 (1991).
	EB	MATSUDA, A., et al., "Nucleosides and Nucleotides. 104. Radical and palladium-catalyzed deoxygenation of the allylic alcohol systems in the sugar moiety of pyrimidine nucleosides," <i>Nucleosides & Nucleotides</i> , 11(2/4):197-226 (1992).
	EC	MIKHAILOV, S.N., et al., "Synthesis and properties of 3'C-methylnucleosides and their phosphoric esters," <i>Carbohydrate Research</i> , 124:75-96 (1983).
	ED	MIKHAILOV, S.N., et al., "Substrate properties of C'-methylnucleoside and C'-methyl-2'-deoxynucleoside 5'-triphosphates in RNA and DNA synthesis reactions catalysed by RNA and DNA polymerases," <i>Nucleosides & Nucleotides</i> , 10(1-3):339-343 (1991).
	EE	MIKHAILOV, S.N., et al., "Hydrolysis of 2'- and 3'-C-methyluridine 2'c3'-cyclic monophosphates and interconversion and dephosphorylation of the resulting 2'- and 3'-monophosphates: Comparison with the reactions of uridine monophosphates," <i>J. Org. Chem.</i> , 57 (15):4122-4126 (1992).
	EF	NUTT, R.F., et al., "Branched-chain sugar nucleosides. III. 3'-C-methyladenine", <i>J.Org. Chem.</i> , 33:1789-1795 (1968).
	EG	OIVANEN, M., et al., "Additional evidence for the exceptional mechanism of the acid-catalyzed hydrolysis of 4-oxopyrimidine nucleosides: Hydrolysis of 1-(1-alkoxyalkyl)uracils, seconucleosides, 3'-C-alkyl nucleosides and nucleoside 3',5'-cyclic monophosphates," <i>J. Chem. Soc. Perkin Trans. 2</i> , 1994:309-314 (1994).
	EH	ONG, S.P., et al., "Synthesis of 3'-C-methyladenosine and 3'-C-methyluridine diphosphates and their interaction with the ribonucleoside diphosphate reductase from <i>Corynebacterium nephridii</i> ," <i>Biochemistry</i> , 31(45):11210-11215 (1992).
	EI	Oral Session V, Hepatitis C Virus, Flaviviridae; 16 th International Conference on Antiviral Research (April 27, 2003, Savannah, Ga.) p A75-77.
	EJ	PAN-ZHOU, X-R., et al., "Differential effects of antiretroviral nucleoside analogs on mitochondrial function in HepG2 cells," <i>Antimicrob. Agents Chemother.</i> , 44:496-503 (2000).
✓	EK	ROSENTHAL, A., et al., "Branched-chain sugar nucleosides. Synthesis of 3'-C-ethyl (and 3'-C-buty1)uridine <i>Carbohydrate Research</i> , 79:235-242 (1980).
(a)	EL	SAMANO, V., et al., "Synthesis and radical-induced ring-opening reactions of 2'-deoxyadenosine-2'-spirocyclopropane and its uridine analogue. Mechanistic probe for ribonucleotide reductases," <i>J. Am. Chem. Soc.</i> , 114:4007-4008 (1992).

Examiner Signature		Date Considered	3/29/04
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**INFORMATION DISCLOSURE
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Sheet 6 of 6				Complete if Known	
				Application Number	10/602,693
				Filing Date	June 20, 2003
				First Named Inventor	Sommadossi <i>et al.</i>
				Group Art Unit	1614
				Examiner Name	Unassigned
				Attorney Docket Number	06171.105070 IDX 1006 DIV

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OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS

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(17)	FA	SAMANO, V., <i>et al.</i> , "Nucleic acid related compounds. 77. 2',3'-Didehydro-2',3'-dideoxy-2'(and 3')-methylnucleosides via [3,3]-sigmatropic rearrangements of 2'(and 3')-methylene-3'(and 2')-O-thiocarbonyl derivatives and radical reduction of a 2'-chloro-3'-methylene analogue," <i>Can. J. Chem.</i> , 71:186-191 (1993).	
	FB	SCHMIT, C., <i>et al.</i> , "The effects of 2'- and 3'-alkyl substituents on oligonucleotide hybridization and stability," <i>Bioorganic & Medicinal Chemistry Letters</i> , 4(16):1969-1974 (1994). ["Altmann"]	
	FC	SERAFINOWSKI, P.J., <i>et al.</i> , "New method for the preparation of some 2'- and 3'-trifluoromethyl-2',3'-dideoxyuridine derivatives," <i>Tetrahedron</i> (Elsevier Science Publishers), 56(2):333-339 (1999).	
	FD	SHARMA, P.K., <i>et al.</i> , "Synthesis of 3'-trifluoromethyl nucleosides as potential antiviral agents," <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 19(4):757-774 (2000).	
	FE	SOMMADODSSI J-P., <i>et al.</i> , "Comparison of cytotoxicity of the (-) and (+)-enantiomer of 2',3'-dideoxy-3'-thiacytidine in normal human bone marrow progenitor cells" <i>Biochemical Pharmacology</i> , 44:1921-1925 (1992).	
	FF	SOMMADODSSI J-P., <i>et al.</i> , "Toxicity of 3'-azido-3'-deoxythymidine and 9-(1,3-dihydroxy-2-propoxymethyl)guanine for normal human hematopoietic progenitor cells in vitro" <i>Antimicrobial Agents and Chemotherapy</i> , 31:452-454 (1987).	
	FG	TRITSCH, D., <i>et al.</i> , "3'-β-ethynyl and 2'-deoxy-3'-β-ethynyl adenosines: First 3'-β-branched adenosines substrates of adenosine deaminase," <i>Bioorganic & Medicinal Chemistry Letters</i> , 10:139-141 (2000).	
	FH	TUNITSKAYA, V.L., <i>et al.</i> , "Substrate properties of C'-methyl UTP derivatives in T7 RNA polymerase reactions. Evidence for N-type NTP conformation," <i>FEBS Letters</i> , 400:263-266 (1997).	
	FI	USUI, H., <i>et al.</i> , "Synthesis of 2'-deoxy-8,2'-ethanoadenosine and 3'-deoxy-8,3'-ethanoadenosine (Nucleosides and Nucleotides. LXIV)," <i>Chem. Pharm. Bull.</i> , 34(1):15-23 (1986).	
	FJ	WALCZAK, K., <i>et al.</i> , "Synthesis of 1-(3-alkyl-2,3-dideoxy-D-pentofuranosyl)uracils with potential anti-HIV activity," <i>Acta Chemica Scand.</i> , 45:930-934 (1991).	
	FK	WALTON, E., <i>et al.</i> , "Branched-chain sugar nucleosides. V. Synthesis and antiviral properties of several branched-chain sugar nucleotides," <i>J. Med. Chem.</i> , 12:306-309 (1969).	
	FL	WOLFE, M.S., <i>et al.</i> , "A concise synthesis of 2'-C-methylribonucleosides," <i>Tetrahedron Letters</i> , 36(42):7611-7614 (1995).	
(3)	FM	WU, J.-C., <i>et al.</i> , "A new stereospecific synthesis of [3.1.0] bicyclic cyclopropano analog of 2',3'-dideoxyuridine," <i>Tetrahedron</i> , 46(7):2587-2592 (1990).	

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